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## IN THE CLAIMS

## Amendments to the Claims

A Listing of Claims is provided as follows and will replace any previous listing. No new matter has been added.

## **Listing of Claims:**

1. (Currently Amended) A method for purifying a target protein from a protein solution containing the target protein by using liquid chromatography, wherein the target protein is glucose dehydrogenase derived from a microorganism belonging to the genus Burkholderia and has  $\alpha$ ,  $\beta$ ,  $\gamma$  subunits, the liquid chromatography comprises comprising:

a first step of introducing the protein solution into a column filled with a packing agent, and causing the packing agent to holding the target protein, the packing agent being an ion-exchange resin containing a quaternary ammonium group as an ion-exchange group; and

a second step of eluting the target protein by using an eluent containing a hydroxy-cholate.

## 2-5. (Canceled)

6. (Currently Amended) The method for purifying protein according to Claim [[3]] 1, wherein the β subunit of the glucose dehydrogenase provides electron transfer protein activity and has a molecular molecular weight of approximately 43 kDa in SDS-gel electrophoresis under a reducing environment, and

the a subunit of the glucose dehydrogenase provides the protein-which has glucose dehydrogenation activity and has having a molecule molecular weight of approximately 60 kDa in SDS-gel electrophoresis under a reducing environment.

7. (Currently Amended) The method for purifying protein according to Claim 1, wherein the hydroxy-cholate comprises a sodium cholate.

8. (Currently Amended) The method for purifying protein according to Claim 1, wherein the hydroxy-cholate in the eluent is maintained at a constant concentration [[in]] during the elution of the target protein from the packing agent.

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- 9. (Original) The method for purifying protein according to Claim 8, wherein the concentration of the hydroxy-cholate in the eluent is selected from a range of 0.5 through 2.5 wt%.
- 10. (Canceled)

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- 11. (Currently Amended) The method for purifying protein according to Claim [[10] 1. wherein the microorganism belonging to the genus Burkholderia is provided by Burkholderia cepacia KS1 strain (FERM BP-7306).
- 12. (Currently Amended) The method for purifying protein according to Claim 3, wherein the glucose dehydrogenase is produced by a transformant,

the transformant being produced by engineering a host microorganism with [[a]] DNA from a microorganism belonging to the genus Burkholderia for coding the electron transfer protein and the protein active against glucose encoding the  $\alpha$  and  $\beta$ . subunits.

- 13. (Currently Amended) The method for purifying protein according to Claim 12, wherein the host microorganism is provided by Pseudomonas putida.
- 14. (Currently Amended) The method for purifying protein according to Claim 12, wherein the host microorganism is provided by E. coli bacterium.
- 15-23. (Canceled)
- 24. (New) The method for purifying protein according to Claim 1, wherein the  $oldsymbol{a}$  and  $oldsymbol{\gamma}$ subunits of the glucose dehydrogenase provide glucose dehydrogenation activity and the

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 $\gamma$  subunit has a molecular weight of approximately 14 kDa in SDS-gel electrophoresis under a reducing environment.